Figure 1 A

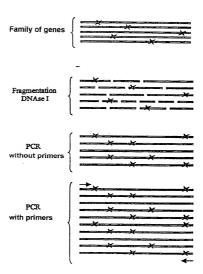


Figure 1 B (Double-stranded process carried out, but illustrated here with a single strand)

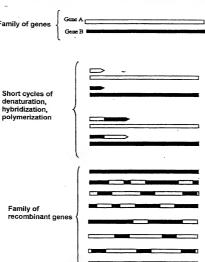
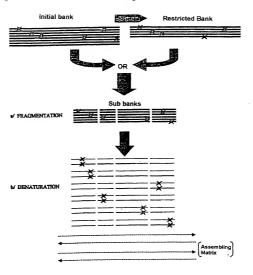


Fig.2



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Fig. 2- continued

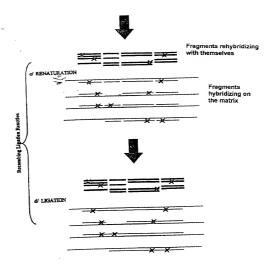
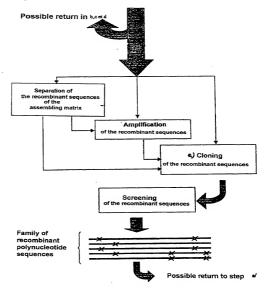


Fig. 2 continued



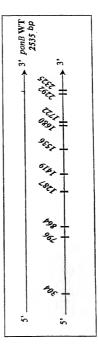


Figure 3: Position of the ten mutation zones (sites Pvull and Pst I)

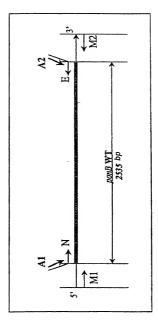


Figure 4: Position of the primers used as compared to the sequence of the son Bgene

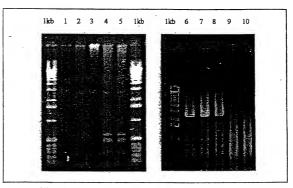


Fig. 5: Migration of the RLR reactions and of the PCR amplifications of these reactions

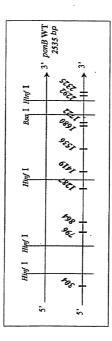


Figure 6: Position of the mutations as compared to the restriction fragments